Kallmann syndrome-heart disease syndrome

INSERM

Source

INSERM. (1999). Orphanet: an online rare disease and orphan drug data base. Kallmann syndrome-heart disease syndrome. ORPHA:2326

Kallmann syndrome with cardiopathy is characterised by hypogonadotropic hypogonadism associated with gonadotropin-releasing hormone (GnRH) deficiency, anosmia or hyposmia (with hypoplasia or aplasia of the olfactory bulbs) and complex congenital cardiac malformations (double-outlet right ventricle, dilated cardiomyopathy, right aortic arch). It represents a distinct clinical entity from Kallmann syndrome.